DEC - 6 2006

Exhibit B 510(K) SUMMARY

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR §807.92(c).

The assigned 510(k) number is: K063500.

1. Submitter:

Shenzhen Mindray Bio-medical Electronics Co., LTD
Mindray Building, Keji 12th Road South, Hi-tech Industrial Park, Nanshan, Shenzhen, 518057, P. R. China

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Contact Person:
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Nanshan, Shenzhen, 518057, P. R. China

Date Prepared: October 16, 2006

2. Device Name: DC-6 Diagnostic Ultrasound System

Classification
Regulatory Class: II
Review Category: Tier II
21 CFR 892.1550 Ultrasonic Pulsed Doppler Imaging System (90-IYN)
21 CFR 892.1560 Ultrasonic Pulsed Echo Imaging System (90-IYO)
21 CFR 892.1570 Diagnostic Ultrasound Transducer (90-ITX)

3. Marketed Device:

DC-6 Diagnostic Ultrasound System is substantially equivalent to the following devices: Toshiba NEMIO SSA-550A (K#010631) and FAMIO SSA-530A (K#051500), Aloka SSD-5000 (K#012080), Philips iU22 (K#042540), Siemens SONOLINE G60 S (K#052894) and Sequoia 512 (K#052410), Mindray DP-9900 (#K061189).

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K063500

4. Device Description:

The DC-6 Diagnostic Ultrasound System is a general purpose, mobile, software controlled, ultrasound diagnostic system. Its function is to acquire and display ultrasound mages in B-Mode, M-Mode, Color mode, PW mode, Power/DirPower mode or the combined mode (i.e. B/M Mode). This system is a Track 3 device that employs an array of probes that include linear array and convex linear array with a frequency range of approximately 2.5 MHz to 12 MHz.

5. Intended Use:

The device is intended for use by a qualified physician for ultrasound evaluation of adult abdomen, adult cardiology, pediatric abdomen, pediatric cardiology, fetal cardiology, gynecology, obstetrics (first trimester, second and third trimester), kidney, prostate, thyroid, breast, and other small parts, and peripheral vascular (peripheral artery and peripheral vein), and carotid scanning.

6. Safety Considerations:

The DC-6 Diagnostic Ultrasound System has been tested as Track 3 Device per the FDA Guidance document "Information for Manufacturers Seeking Marketing Clearance of Diagnostic Ultrasound Systems and Transducers" issued September 1997. The acoustic output is measured and calculated per NEMA UD 2 Acoustic Output Measurement Standard for Diagnostic Ultrasound Equipment: 2004 and NEMA UD 3 Output Display Standard. The device conforms to applicable medical device safety standards, such as IEC 60601-1, IEC 60601-1-2, IEC 60601-2-37 and ISO 10993-1.

Conclusion:

The conclusions drawn from testing of the DC-6 Diagnostic Ultrasound System demonstrate that the device is as safe and effective as the legally marketed predicate devices.

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DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

DEC 14 2006

Shenzhen Mindray Bio-Medical Electronics Co., Ltd.
% Mr. Robert Mosenkis
President
Citech
5200 Butler Pike
PLYMOUTH MEETING PA 19462-1298

Re: K063500
Trade Name: DC-6 Diagnostic Ultrasound System
Regulation Number: 21 CFR 892.1550
Regulation Name: Ultrasonic pulsed doppler imaging system
Regulation Number: 21 CFR 892.1560
Regulation Name: Ultrasonic pulsed echo imaging system
Regulation Number: 21 CFR 892.1570
Regulation Name: Diagnostic ultrasonic transducer
Regulatory Class: II
Product Code: IYN, IYO, and ITX
Dated: November 17, 2006
Received: November 20, 2006

Dear Mr. Mosenkis:

This letter corrects our substantially equivalent letter of December 6, 2006.

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

This determination of substantial equivalence applies to the following transducers intended for use with the DC-6 Diagnostic Ultrasound System, as described in your premarket notification:

Transducer Model Number			
3C5	3C1	6CV1	7L4

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus permits your device to proceed to market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html

If you have any questions regarding the content of this letter, please contact Ralph Shuping at (240) 276-3666.

Sincerely yours,

for Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure(s)

Diagnostic Ultrasound Indications for Use Form

System × Transducer _____

Model: DC-6

510(k) Number(s) K063500

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other* (specify)
Ophthalmic										
Fetal		N	N	N		N	N		N	N
Abdominal		N	N	N		N	N		N	N
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric		N	N	N		N	N		N	N
Small organ(specify)**		N	N	N		N	N		N	
Neonatal Cephalic		N	N	N		N	N		N	
Adult Cephalic										
Cardiac		N	N	N		N	N		N	N
Transesophageal										
Transrectal		N	N	N		N	N		N	
Transvaginal		N	N	N		N	N		N	
Transurethral										
Intravascular										
Peripheral Vascular		N	N	N		N	N		N	
Laparoscopic										
Musculo-skeletal Conventional		N	N	N		N	N		N	
Musculo-skeletal Superficial		N	N	N		N	N		N	
Other (specify)										

N=new indication; P=previously cleared by FDA; E=added under Appendix E

Additional comments: Combined modes: B+M, PW+B, Color + B, Power + B, PW +Color+ B, Power + PW +B.

*Other: Tissue Harmonic Imaging. The feature does not use contrast agents.

**Small organ-breast, thyroid, testes, etc.

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Concurrence of CDRH, Office of Device Evaluation(ODE)

David A. Lyman 0047

(Division Sign-Off)

Division of Reproductive, Abdominal, and Radiological Devices

510(k) Number K063500

Prescription USE (Per 21 CFR 801.109)

Diagnostic Ultrasound Indications for Use Form

System _____ Transducer X

Model: 3C5

510(k) Number(s) K063500

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other* (specify)
Ophthalmic										
Fetal		N	N	N		N	N		N	N
Abdominal		N	N	N		N	N		N	N
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small organ(specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N=new indication; P=previously cleared by FDA; E=added under Appendix E

Additional comments: Combined modes: B+M, PW+B, Color + B, Power + B, PW +Color+ B, Power + PW +B.

*Other: Tissue Harmonic Imaging. The feature does not use contrast agents.

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Concurrence of CDRH, Office of Device Evaluation(ODE)

David A. Legson 0048

(Division Sign-Off)
Division of Reproductive, Abdominal, and Radiological Devices
510(k) Number K063500

Prescription USE (Per 21 CFR 801.109)

Diagnostic Ultrasound Indications for Use Form

System _____ Transducer X

Model: 3C1

510(k) Number(s) K063500

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other* (specify)
Ophthalmic										
Fetal										
Abdominal		N	N	N		N	N		N	N
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric		N	N	N		N	N		N	N
Small organ(specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac		N	N	N		N	N		N	N
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N=new indication; P=previously cleared by FDA; E=added under Appendix E

Additional comments: Combined modes: B+M, PW+B, Color + B, Power + B, PW+Color+ B, Power + PW +B.

*Other: Tissue Harmonic Imaging. The feature does not use contrast agents.

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Concurrence of CDRH, Office of Device Evaluation(ODE)

David A. Lagerwall 0643

(Division Sign-Off)
Division of Reproductive, Abdominal, and Radiological Devices
510(k) Number K063500

Prescription USE (Per 21 CFR 801.109)

Diagnostic Ultrasound Indications for Use Form

System ____ Transducer ×

Model: 6CV1

510(k) Number(s) K063500

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small organ(specify)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		N	N	N		N	N		N	
Transvaginal		N	N	N		N	N		N	
Transurethral										
Intravascular										
Peripheral Vascular										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N=new indication; P=previously cleared by FDA; E=added under Appendix E

Additional comments: Combined modes: B+M, PW+B, Color + B, Power + B, PW +Color+ B, Power + PW +B.

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Concurrence of CDRH, Office of Device Evaluation(ODE)

David A. Lyerman 0050

(Division Sign-Off)

Division of Reproductive, Abdominal, and Radiological Devices

510(k) Number K063500

Prescription USE (Per 21 CFR 801.109)

Diagnostic Ultrasound Indications for Use Form

System _____ Transducer X

Model: 7L4

510(k) Number(s) K063500

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (specify)	Other (specify)
Ophthalmic										
Fetal										
Abdominal										
Intraoperative (specify)										
Intraoperative Neurological										
Pediatric										
Small organ(specify)**		N	N	N		N	N		N	
Neonatal Cephalic		N	N	N		N	N		N	
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral Vascular		N	N	N		N	N		N	
Laparoscopic										
Musculo-skeletal Conventional		N	N	N		N	N		N	
Musculo-skeletal Superficial		N	N	N		N	N		N	
Other (specify)										

N=new indication; P=previously cleared by FDA; E=added under Appendix E

Additional comments: Combined modes: B+M, PW+B, Color + B, Power + B, PW +Color+ B, Power + PW +B.

**Small organ-breast, thyroid, testes, etc.

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Concurrence of CDRH, Office of Device Evaluation(ODE)

Donald B. Seyman 0051

(Division Sign-Off)

Division of Reproductive, Abdominal, and Radiological Devices

510(k) Number K063500

Prescription USE (Per 21 CFR 801.109)